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Application No.	10/821667	Prepared by	the	Tracking Number	0587 2434
Examiner-GAU	Sotola - 1626	Date	11/14/04	Week Date	12/8/03
		No. of queries	2	CWU (JFW)	

JACKET			
a. Serial No.	f. Foreign Priority	k. Print Claim(s)	p. PTO-1449
b. Applicant(s)	g. Disclaimer	l. Print Fig.	q. PTOL-85b
c. Continuing Data	h. Microfiche Appendix	m. Searched Column	r. Abstract
d. PCT	i. Title	n. PTO-270/328	s. Sheets/Figs
e. Domestic Priority	j. Claims Allowed	o. PTO-892	t. Other

SPECIFICATION

a. Page Missing
b. Text Continuity
c. Holes through Data
d. Other Missing Text
e. Illegible Text
f. Duplicate Text
g. Brief Description
h. Sequence Listing
i. Appendix
j. Amendments
(k.) Other

CLAIMS

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b. Improper Dependency
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d. Incorrect Numbering
e. Index Disagrees
f. Punctuation
g. Amendments
h. Bracketing
i. Missing Text
j. Duplicate Text
k. Other

MESSAGE ① Chemical structure under EXAMPLE 6 has handwritten date and a box drawn, should this be picked up?

② The illustrations on pages 56 and 69 do not come within the exceptions of 37CFR 1.58(a). Please furnish drawings per 37CFR 1.81, delete illustrations from specification and amend text references as necessary.

Thank you.

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RESPONSE ① No. Clean copy of page attached

② The illustrations are chemical formulas and do not need to be made into figures. 1.58(a) is not applicable to these.

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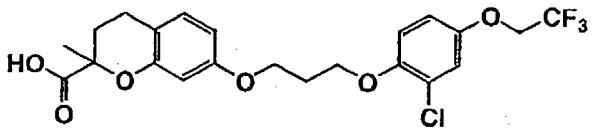
was extracted twice with AcOEt. The combined organic layers were dried over anhydrous Na_2SO_4, filtered, and concentrated to give the title compound 16mg.

^{1}H NMR (500 MHz, $CDCl_3$): δ 7.038 (d, 1H, J=3Hz), 6.979 (d, 1H, J=8.2 Hz), 6.92 (d, 1H, J=9.2 Hz), 6.832 (dd, 1H, J=3, J=9.2 Hz), 6.547 (m, 2H), 4.73 (dd, 1H), 4.313 (q, 2H), 4.188 (t, 4H), 2.741-2.868 (m, 2H), 2.40 (m, 1H), 2.29 (p, 2H), 2.163 (m, 1H). ms: m/e=461 (M+1).

EXAMPLE 6

7-(3-(2-Chloro-4-(2,2,2-trifluoroethoxy)phenoxy)propoxy)-2-methylchromane-2-carboxylic acid



The title compound was prepared from ethyl 7-(3-(2-chloro-4-(2,2,2-trifluoroethoxy)phenoxy)propoxy)-chromane-2-carboxylate (Example 5, Step D) following the procedure described in Example 1, Step C employing iodomethane instead of iodoethane followed by hydrolysis as described in Example 5, Step E.

^{1}H NMR (500 MHz, $CDCl_3$): δ 7.032 (d, 1H, J =3.0 Hz), 6.952 (d, 1H), 6.916 (d, 1H, J=8.9 Hz), 6.836 (dd, 1H, J=3.0, 8.9 Hz), 6.514 (m, 2H), 4.31 (q, 2H, J=8 Hz), 4.177 (m, 4H), 2.718 (m, 2H), 2.389 (dt, 1H, J=5.0 Hz, 13.7 Hz), 2.285 (pent, 2H, J=5.9 Hz), 1.953 (dt, 1H, J=8.2 Hz, 13.5 Hz), 1.661 (s, 3H). ms: m/e=475 (M+1).

EXAMPLE 7

7-(3-(2-Chloro-4-(2,2,2-trifluoroethoxy)phenoxy)propoxy)-2-ethylchromane-2-carboxylic acid